RSC Mechanochemistry



PAPER

View Article Online
View Journal | View Issue



Cite this: RSC Mechanochem., 2024, 1,

Direct arylation of alkyl fluorides using in situ mechanochemically generated calcium-based heavy Grignard reagents†

Pan Gao,^a Julong Jiang,^b Yamato Fukuzawa,^a Satoshi Maeda, ^b ^c Koji Kubota ^b *ac and Hajime Ito *b *ac

Here, we report the reaction of calcium-based heavy Grignard reagents, which are easily generated by a mechanochemical method, with unactivated alkyl fluorides in the absence of transition metal catalysts to produce the corresponding arylated products in moderate to good yields. This is the first example of the nucleophilic substitution of an inert $C(sp^3)$ -F bond by an organocalcium species. Preliminary mechanistic studies based on theoretical calculations indicate that tetrameric aryl calcium species facilitate the unprecedented $C(sp^3)$ -F bond arylation.

Received 21st June 2024 Accepted 13th August 2024

DOI: 10.1039/d4mr00067f

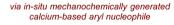
rsc.li/RSCMechanochem

Organocalcium reagents have demonstrated remarkable synthetic potential as carbon nucleophiles because they frequently exhibit distinct reactivity owing to the large electronegativity difference between calcium and carbon. However, in contrast to the extensive research on traditional organometallic carbon nucleophiles, such as Grignard reagents, studies on synthetic applications involving calcium-based nucleophiles are scarce. This is due to a lack of operationally simple protocols for obtaining these nucleophiles from commercially available starting materials, which may impede comprehensive investigations into the reactivity profile of calcium-based carbon nucleophiles; this in turn can result in basic reactions being overlooked. Legal 2015

We recently developed a simple mechanochemical protocol using ball milling to generate calcium-based heavy Grignard reagents (R-CaX)⁷⁻¹³ from commercially available, unactivated calcium metal. ¹⁴ Conventionally, such direct synthesis necessitates the production of activated calcium metal (Rieke calcium) under harsh conditions using alkali metals. ¹⁰⁻¹² However, in our mechanochemical approach, the strong mechanical impact during ball milling may provide sufficient energy to overcome the high atomization energy of calcium metal, enabling the direct generation of calcium-based nucleophiles from commercial calcium metal without the need for laborious preactivation processes. ^{14,15} Furthermore, all synthetic operations

can be carried out under ambient conditions. We discovered a new reactivity of aryl calcium nucleophiles using this operationally simple protocol to rapidly generate organocalcium nucleophiles; they reacted smoothly with alkyl iodides or bromides, yielding the corresponding arylated products in good-to-high yields.¹⁴

In this study, we discovered that direct arylation of alkyl fluorides using *in situ* mechanochemically generated aryl calcium nucleophiles is highly efficient even in the absence of transition metal catalysts (Scheme 1).^{16,17} This is the first example of the nucleophilic substitution of an inert C(sp³)–F bond by an organocalcium species.^{18,19} Although direct C(sp³)–F bond arylation is also feasible using Grignard reagents, as discovered by Matsubara *et al.*,^{16a} the present study sheds light on the previously overlooked reactivity of calcium-based heavy Grignard reagents. The developed organocalcium-mediated arylation exhibited a significantly shorter reaction time (1 h) than that under Matsubara's conditions using Grignard reagents (>12 h),^{16a} which is likely owing to the higher



- The first nucleophilic substitution of C(sp³)-F bond by organocalcium
- Simple mechanochemical protocol for rapid generation of organocalcium
- Preliminary theoretical calculations to reveal the unprecedented reactivity

Scheme 1 Arylation of alkyl fluorides using mechanochemically generated calcium-based heavy Grignard reagents.

[&]quot;Division of Applied Chemistry, Faculty of Engineering, Hokkaido University, Sapporo, Hokkaido, Japan. E-mail: kbt@eng.hokudai.ac.jp; hajito@eng.hokudai.ac.jp

^bDepartment of Chemistry, Faculty of Science, Hokkaido University, Sapporo, Hokkaido, Japan

^cInstitute for Chemical Reaction Design and Discovery (WPI-ICReDD), Hokkaido University, Sapporo, Hokkaido, Japan

[†] Electronic supplementary information (ESI) available. See DOI: https://doi.org/10.1039/d4mr00067f

nucleophilicity of the organocalcium species. Density functional theory (DFT) calculations were used to propose the mechanism of the C(sp3)-F bond arylation. This study demonstrates the effectiveness of a simple mechanochemical protocol to explore the novel reactivity of organocalcium species more easily.

All the reactions were conducted in a Retsch MM400 mill (stainless-steel milling jar; 30 Hz; stainless-steel balls). Based on our previous study,14 the reaction between iodobenzene (1a, 3.0 equiv.) and 1-fluorododecane (2a, 1.0 equiv.) was performed in the presence of commercially available calcium metal (2.0 equiv.) and tetrahydropyran (THP) (4.0 equiv.) in a 1.5 mL stainless steel jar with two stainless steel balls (diameter: 7 mm) in air. Following 60 min of ball milling (30 Hz) using a heat gun as a temperature controller²⁰ to maintain an internal temperature of 80 °C (for details, refer to the ESI†), the C(sp³)-F bond arylation product 3aa was successfully obtained in good yield (86%, Table 1, entry 1). When the reaction was performed at room temperature without the use of a heat gun, the yield of 3aa decreased significantly (28%; Table 1, entry 2). Ether additives are thought to coordinate with the calcium(II) center to form thermodynamically stable species,5 facilitating the formation of aryl calcium nucleophiles under mechanochemical conditions.14 The reaction did not proceed without the addition of THP (<5%; Table 1, entry 3). The use of tetrahydrofuran (THF) resulted in a moderate yield (51%; Table 1, entry 4). The reaction was completely suppressed when dimethoxyethane (DME)

was used (<5%; Table 1, entry 5). The reaction with two equivalents of THP gave a slightly lower yield of 3aa (71%, Table 1, entry 6). Lowering the vibration frequency of ball milling resulted in a significant decrease in yield, indicating that mechanical impact played an important role in this transformation (62%; Table 1, entry 7). Lower yields were obtained when the amounts of 1a and calcium metal were reduced (61 and 41%, respectively; Table 1, entries 8 and 9). When the reaction was performed in a 10 mL stainless-steel jar with two stainless-steel balls (diameter: 10 mm), a moderate yield of 3aa was obtained (66%; Table 1, entry 10). Given the possibility of cross-electrophile coupling catalyzed by transition metals derived from stainless-steel milling jars and/or balls,17 we performed the reaction in a commercially available 10 mL zirconium oxide jar with two zirconium oxide balls (diameter: 10 mm), yielding the desired arylation product 3aa (43%; Table 1, entry 11). This result suggests that the C(sp3)-F bond arylation process is likely to proceed only by organocalcium species without catalysis of transition metal contamination.¹⁷ We believe that the lower mechanical impact of zirconium oxide balls, which are much lighter than stainless-steel balls, is a primary factor contributing to the observed decrease in yield. The mechanical activation of calcium metal is crucial for the formation of organocalcium compounds, and the reduced impact with zirconium oxide balls likely hampers this activation process. While Matsubara et al. reported that phenyl Grignard reagent reacted with alkyl fluorides in THF solution, 16a the use

Optimization of the cross-electrophile coupling between iodobenzene (1a) and 1-fluorododecane (2a) in the presence of calcium Table 1 metala

Entry	Deviation from "standard conditions"	Yield (%) of 3aa ^b
1	None	86 (74)
2	Room temperature	28
3	Without THP	<5
4	THF instead of THP	51
5	DME instead of THP	<5
6	2.0 equiv. of THP	71
7	Frequency: 15 Hz	62
8	2.0 equiv. of 1a	61
9	1.0 equiv. of Ca metal	41
10	10 mL jar with a 10 mm ball \times 2 (stainless-steel)	66
11	10 mL jar with 10 mm ball \times 2 (zirconium oxide)	43
12	Mg metal instead of Ca metal	<5
13	In THP solution (0.2 M) using a test tube	<5
14	Bromobenzene instead of iodobenzene (1a)	<5
15	Chlorobenzene instead of iodobenzene (1a)	<5

^a Conditions: 1a (1.5 mmol), 2a (0.5 mmol), Ca (0.75 mmol), and THP (2.0 mmol) in a stainless-steel ball-milling jar (1.5 mL) with two stainless-steel balls (diameter: 7 mm). b Yields determined by HNMR spectroscopy using dibromo methane as an internal standard. The isolated yields are shown in parentheses.

of magnesium instead of calcium did not result in the formation of arylation product 3aa under mechanochemical conditions (<5%; Table 1, entry 12).²¹ To verify the role of the ballmilling technique, we performed the reaction in THP solution at 80 °C (Table 1, entry 13). We discovered that 3aa was not detected, and almost no conversion of 1a was observed (3%), suggesting that the mechanical impact of ball milling is required for the generation of the aryl calcium nucleophile, which subsequently reacts with alkyl fluoride 2a. Because it is difficult to generate the corresponding calcium nucleophiles, using bromo- or chlorobenzene instead of 1a resulted in no product formation (Table 1, entries 14 and 15).

Using the obtained optimal conditions, the substrate scope was investigated. We first examined aryl iodides bearing different substituents (Table 2). The introduction of substituents such as methyl (1b), *tert*-butyl (1c), phenyl (1d), or diphenylamino (1e) groups at the *para*-position of the phenyl ring exhibited no effect on reactivity, resulting in the desired arylation products (3ba–3ea) in moderate to good yields (62–83%). Substrates with a methyl group at the meta position (1f) also reacted under mechanochemical conditions to produce the

Table 2 Substrate scope for aryl iodides^{a,b}

target product (3fa) in high yield (88%). The reaction with a substrate containing a sterically demanding substituent, such as the 2-isopropyl group (1g), proceeded smoothly, producing the corresponding product (3ga) in high yield (80%). We also discovered that arylation using polycyclic aryl iodides (1h and 1i) was feasible and produced the desired products (3ha and 3ia) in high yields (69 and 85%, respectively). However, the reactions failed when functional groups, including methoxy (1j), ester (1k), or carbonyl (1l) groups, were present.

We then investigated the substrate scope in terms of alkyl fluorides (Table 3). Alkyl fluoride with a phenyl group 2b reacted smoothly to produce the desired product 3ab in good yield (82%). The presence of a trifluoromethyl group on the benzene ring (2c) had no effect on the reaction, and only the C-F bond of the monofluoromethyl group located at the end of the alkyl chain was activated to produce the corresponding arylation product 3ac in high yield (86%). Under the optimized conditions, functional groups, including phenoxy (2d), phenyl thioether (2e), and alkenes (2f and 2g), were tolerated, yielding the target products (3ad-3ag) in moderate to good yields (70-86%). However, it is important to note the current limitations of this method, including the difficulty of coupling with alkyl fluorides containing a benzyloxy group (2h) or a thiophene moiety (2i), resulting in low yields of the desired products (3ah and 3ai in 21 and 20% yields, respectively). Secondary alkyl fluoride 2j was used; however, the corresponding arylation product was not detected.

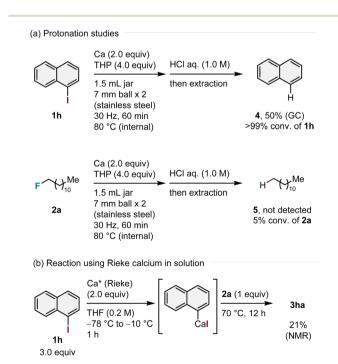
 Table 3
 Substrate scope with respect to alkyl fluorides a,b

 $[^]a$ Conditions: 1 (1.5 mmol), 2a (0.5 mmol), Ca (0.75 mmol), THP (2.0 mmol), in a stainless-steel ball-milling jar (1.5 mL) with two stainless-steel balls (diameter: 7 mm). b Yields determined by 1 H NMR spectroscopy using dibromo methane as an internal standard. The isolated yields are shown in parentheses.

^a Conditions: 1a (1.5 mmol), 2 (0.5 mmol), Ca (0.75 mmol), and THP (2.0 mmol) in a stainless-steel ball-milling jar (1.5 mL) with two stainless-steel balls (diameter: 7 mm). ^b Yields determined by ¹H NMR spectroscopy using dibromomethane as an internal standard. The isolated yields are shown in parentheses.

Control experiments were performed to obtain mechanistic insights (Scheme 2). When iodonaphthalene (1h) was subjected to standard conditions in the absence of alkyl fluorides, protonation product 4 was obtained in 50% yield after quenching with 1.0 M aq. HCl, indicating that aryl calcium species formed following 1h under mechanochemical conditions (Scheme 2a).14 In contrast, the reaction using alkyl fluoride 2a in the absence of 1h yielded no protonation product 5, and 2a converted at a low rate (5% conversion, Scheme 2a). On the basis of protonation studies, we propose that C(sp³)-F bond arylation occurs most likely through the selective formation of aryl calcium nucleophiles via direct insertion of calcium metal into the C(sp²)-I bond, followed by nucleophilic substitution of alkyl fluorides.14 To further support our hypothesis, we prepared an arvl calcium reagent using pre-synthesized reagent, Rieke calcium, in solution and subjected it to nucleophilic substitution (Scheme 2b).10-12 We discovered that the prepared aryl calcium reagent reacted with 1-fluorododecane (2a) to produce the desired product 3ha (21%, Scheme 2b), supporting our hypothesis.

Furthermore, to gain insight into the mechanism of nucleophilic substitution, DFT calculations were performed on the model substrates, iodobenzene (1a) and n-butyl fluoride. Preliminary calculations indicated that the tetracoordinate mononuclear calcium complex was highly unstable owing to the presence of vacant coordination sites on the calcium atom. Meanwhile, the artificial force-induced reaction (AFIR)-based sampling calculation²² using the tetrameric model yielded 1787 possible structures, the most stable of which exhibits a highly symmetric cubic structure (Fig. 1a).23 The coordination vacancies of each calcium atom in this cubic structure are



Scheme 2 Control experiments.^a Details of the reaction conditions are provided in the ESI.†

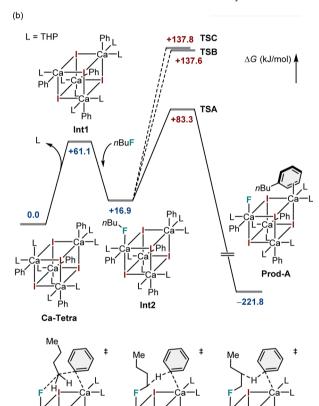


Fig. 1 Theoretical calculations to reveal the plausible structure of PhCal(THP)₂ in solid state and the mechanism of the defluorinated C-C cross-coupling reaction (energies are given in kJ mol^{-1}). (a) Stablity of calcium species. (b) Energy profiles of possible reaction pathways

TSB

fulfilled, resulting in a coordination number of six. Furthermore, DFT calculations demonstrated that the tetramer was more stable than the dimer (Fig. 1a). Therefore, the tetramer with the composition formula (PhCaI)(THP)₂ is proposed as the most likely structure in the reaction system.

After confirming the possible structure of the aryl calcium nucleophile, we focused on elucidating the detailed mechanism

TSC

of defluoroarylation. Fig. 1b depicts three possible reaction pathways, beginning with the tetrameric species Ca-Tetra. To generate a vacant site for alkyl fluoride coordination, one of the THP ligands must be left in the first instance. This step increases free energy by $61.1 \text{ kJ} \text{ mol}^{-1}$. The alkyl fluoride substrate binds to the five-coordinate calcium atom through its fluorine atom, lowering the free energy by 44.2 kJ mol⁻¹. Following the formation of Int2, three possible reaction pathways emerge: arylation (TSA), alpha-H abstraction reaction (TSB), and beta-H abstraction reaction (TSC). The arylation reaction has the lowest barrier among the three possible pathways, with an overall free energy barrier of +83.3 kJ mol⁻¹. The calculated results are consistent with our experimental findings. In this transition state, the initial binding of the fluorine atom to the calcium atom significantly weakens the $C(sp^3)$ -F bond, as calcium(II) is regarded as a moderately strong Lewis acid for the fluorine atom.24 At the transition state, the phenyl anion from another calcium center directly attacks the alphacarbon of the alkyl fluoride, while the previously mentioned calcium atom accepts the fluorine atom as a leaving group. The combination of these two effects causes the chemically inert C(sp³)-F bond to be cleaved, resulting in the formation of a new $C(sp^3)-C(sp^2)$ bond. The ESI† incorporates the visualized geometries of the intermediate and transition states shown in Fig. 1.

In conclusion, we discovered a new reaction of calciumbased heavy Grignard reagents, which were generated in situ via a simple mechanochemical method, with unactivated alkyl fluorides in the absence of transition-metal catalysts, resulting in moderate to high yields of the corresponding C(sp³)-F bond arylation products. Notably, this is the first reported instance of an organocalcium species engaging in nucleophilic substitution of an inert C(sp³)-F bond. To elucidate the unprecedented reactivity of organocalcium nucleophiles in activating inert C(sp³)-F bonds, preliminary theoretical calculations were performed. The findings of this study validate that our operationally simple mechanochemical protocol for rapidly generating organocalcium nucleophiles using commercial, inactivated calcium metal provides a platform for discovering new reactions that can be performed using only calcium-based carbon nucleophiles.

Data availability

The data supporting this article have been included as part of the ESI. \dagger

Conflicts of interest

There are no conflicts to declare.

Acknowledgements

This work was financially supported by the Japan Society for the Promotion of Science (JSPS) *via* KAKENHI grants 22H00318, 22K18333, 24H00453, 24H01832 and 24H01050, by the JST *via* CREST grant JPMJCR19R1, by FOREST grant JPMJFR201I, and

by the Institute for Chemical Reaction Design and Discovery (ICReDD) established by the World Premier International Research Initiative (WPI), MEXT, Japan. We thank Mr Mahiro Takahashi for his help in cross-checking experiments.

Notes and references

- For selected reviews on organocalcium chemistry, see(a)
 P. Hanusa, *Chem. Rev.*, 1993, 93, 1023; (b)
 M. Westerhausen, *Angew. Chem., Int. Ed.*, 2001, 40, 2975; (c)
 J. S. Alexander, *Eur. J. Inorg. Chem.*, 2002, 2002, 2761.
- 2 Selected examples and reviews of the reactions using organocalcium compounds as catalysts:(a) S. Harder, *Chem. Rev.*, 2010, **110**, 3852; (b) S. Kobayashi and Y. Yamashita, *Acc. Chem. Res.*, 2011, **44**, 58; (c) M. S. Hill, D. J. Liptrota and C. Weetman, *Chem. Soc. Rev.*, 2016, **45**, 972; (d) M. Magre, S. Marcin and M. Rueping, *Chem. Rev.*, 2022, **122**, 8261; (e) L. Zhao, P. Deng, X. Gong, X. Kang and J. Cheng, *ACS Catal.*, 2022, **12**, 7877.
- 3 Selected examples of reactions using organocalcium compounds as carbon nucleophiles:(a) A. S. Wilson, M. S. Hill, M. F. Mahon, C. Dinoi and L. Maron, *Science*, 2017, 358, 1168; (b) B. M. Wolf, C. Stuhl, C. Maichle-Mössmer and R. Anwander, *J. Am. Chem. Soc.*, 2018, 140, 2373; (c) K. G. Pearce, C. Dinoi, M. S. Hill, M. F. Mahon, L. Maron, R. S. Schwamm and A. S. S. Wilson, *Angew. Chem., Int. Ed.*, 2022, 61, e202200305.
- 4 Selected examples of organocalcium-mediated polymerization are as follows: (a) S. Harder, F. Feil and A. Weeber, *Organometallics*, 2001, 20, 1044; (b) S. Harder and F. Feil, *Organometallics*, 2002, 21, 2268; (c) M. Westerhausen, S. Schneiderbauer, A. N. Kneifel, Y. Söltl, P. Mayer, H. Nöth, Z. Zhong, P. J. Dijkstra and J. Feijen, *Eur. J. Inorg. Chem.*, 2003, 2003, 3432; (d) Y. Li, H. Deng, W. Brittain and M. S. Chisholm, *Polym. Bull.*, 1999, 42, 635; (e) M.-W. Hsiao and C.-C. Lin, *Dalton Trans.*, 2013, 42, 2041.
- 5 Selected reviews of calcium-based heavy Grignard reagents: (a)
 M. Westerhausen, M. Gärtner, R. Fischer, J. Langer, L. Yu and
 M. Reiher, Chem.-Eur. J., 2007, 13, 6292; (b) M. Westerhausen,
 M. Gärtner, R. Fischer and J. Langer, Angew. Chem., Int. Ed.,
 2007, 46, 1950; (c) M. Westerhausen, Coord. Chem. Rev.,
 2008, 252, 1516; (d) M. Westerhausen, A. Koch, H. Görls and
 S. Krieck, Chem.-Eur. J., 2017, 23, 1456; (e) A. Koch,
 Q. Dufrois, M. Wirgenings, H. Görls, S. Krieck, M. Etienne,
 G. Pohnert and M. Westerhausen, Chem.-Eur. J., 2018, 24,
 16840; (f) S. Harder and J. Langer, Nat. Rev. Chem., 2023, 7,
 843; (g) P. Schüler, S. Sengupta, S. Krieck and
 M. Westerhausen, Chem.-Eur. J., 2023, 29, e202300833.
- 6 (a) Organometallics in Synthesis: A Manual, ed. M. Schlosser, Wiley, Chester, 2nd edn, 2013; (b) T. Banno, Y. Hayakawa and M. Umeno, J. Organomet. Chem., 2002, 653, 288; (c) Handbook of Grignard Reagents, ed. G. S. Silverman and P. E. Rakita, Marcel Dekker, 1996; (d) The Chemistry of Organozinc Compounds, ed. Z. Rappoport and I. Marek, John Wiley & Sons, Chichester, U.K., 2006; (e) U. Wietelmann and J. Klett, Z. Anorg. Allg. Chem., 2018, 644, 194.

- 7 (a) M. Gärtner, H. Görls and M. Westerhausen, Organometallics, 2007, 26, 1077; (b) M. Gärtner, H. Görls and M. Westerhausen, J. Organomet. Chem., 2008, 693, 221; (c) M. Westerhausen, Angew. Chem., Int. Ed., 2009, 48, 5741; (d) J. Langer, M. Köhler, H. Görls and M. Westerhausen, Chem.-Eur. J., 2014, 20, 3154.
- 8 (a) R. Fischer, M. Gärtner, H. Görls, L. Yu, M. Reiher and M. Westerhausen, Angew. Chem., Int. Ed., 2007, 46, 1618; (b) P. Schüler, S. Sengupta, A. Koch, H. Görls, S. Krieck and M. Westerhausen, Chem.-Eur. J., 2022, 28, e202201897; (c) P. Schüler, S. Sengupta, S. Krieck and M. Westerhausen, Chem.-Eur. J., 2023, 29, e20230083.
- 9 A. Koch, M. Wirgenings, S. Krieck, H. Görls, G. Pohnert and M. Westerhausen, Organometallics, 2017, 36, 3981.
- 10 (a) T.-C. Wu, H. Xiong and R. D. Rieke, *J. Org. Chem.*, 1990, 55, 5045; (b) K. Mochida and H. Ogawa, J. Organomet. Chem., 1983, 243, 131; (c) R. D. Rieke, T.-C. Wu and L. I. Rieke, Org. Synth., 1995, 72, 147; (d) D. Bryce-Smith and A. C. Skinner, J. Chem. Soc., 1963, 577; (e) M. L. Hays and T. P. Hanusa, Tetrahedron Lett., 1995, 36, 2435.
- 11 J. Langer, M. Köhler, H. Görls and M. Westerhausen, J. Organomet. Chem., 2014, 751, 563.
- 12 (a) R. Fischer, M. Gärtner, H. Görls and M. Westerhausen, Organometallics, 2006, 25, 3496; (b) M. Gärtner, H. Görls and M. Westerhausen, Synthesis, 2007, 5, 725; (c) M. Westerhausen, M. Gärtner, R. Fischer and J. Langer, Angew. Chem., Int. Ed., 2007, 46, 1950; (d) N. Kawabata, A. Matsumura and S. Yamashita, Tetrahedron, 1973, 29,
- 13 (a) M. Köhler, J. Langer, H. Görls and M. Westerhausen, Organometallics, 2014, 33, 6381; (b) M. Gärtner, H. Görls and M. Westerhausen, J. Organomet. Chem., 2008, 693, 221.
- 14 P. Gao, J. Jiang, S. Maeda, K. Kubota and H. Ito, Angew. Chem., Int. Ed., 2022, 61, e202207118.
- 15 For a tutorial review on the mechanical activation of zerovalent metals, A. C. Jones, J. A. Leitch, S. E. Raby-Buck and D. L. Browne, Nat. Synth., 2022, 1, 763.
- 16 Selected examples of transition-metal-free nucleophilic substitution of alkyl fluorides using organometallic reagents: (a) K. Matsubara, T. Ishibashi and Y. Koga, Org. Lett., 2009, 11, 1765; (b) J. Terao, S. A. Begum, Y. Shinohara, M. Tomita, Y. Naitoh and N. Kambe, Chem. Commun., 2007, 855; (c) L. Kane, B. C. Figula,

- K. Balaraman, J. A. Bertke and C. Wolf, Nat. Commun., 2024, 15, 1866,
- 17 Selected examples and reviews of transition metal catalysedcross-coupling reactions between alkyl fluorides and organometallic reagents: (a) J. Terao, A. Ikumi, H. Kuniyasu and N. Kambe, J. Am. Chem. Soc., 2003, 125, 5646; (b) J. Terao, H. Todo, H. Watanabe, A. Ikumi and N. Kambe, Angew. Chem., Int. Ed., 2004, 43, 6180; (c) J. Terao, H. Watabe and N. Kambe, J. Am. Chem. Soc., 2005, 127, 3656; (d) J. Terao and N. Kambe, Acc. Chem. Res., 2008, 41, 11545; (e) Z. Mo, Q. Zhang and L. Deng, Organometallics, 2012, 31, 6518; (f) L. W. Erickson, E. L. Lucas, E. J. Tollefson and E. R. Jarvo, J. Am. Chem. Soc., 2016, 138, 14006; (g) T. Iwasaki, K. Yamashita, H. Kuniyasu and N. Kambe, Org. Lett., 2017, 19, 3691; (h) T. Hatakeyama, S. Ito, M. Nakamura and E. Nakamura, J. Am. Chem. Soc., 2005, 127, 14192.
- 18 For selected reviews on C-F bond activation in organic synthesis, see: (a) H. Amii and K. Uneyama, Chem. Rev., 2009, **109**, 2119; (b) J.-D. Hamel and J.-F. Paquin, Chem. Commun., 2018, 54, 10224; (c) T. Stahl, H. F. T. Klare and M. Oestreich, ACS Catal., 2013, 3, 1578; (d) G. Coates, F. Rekhroukh and M. R. Crimmin, Synlett, 2019, 30, 2233; (e) T. Ahrens, J. Kohlmann, M. Ahrens and T. Braun, Chem. Rev., 2015, 115, 931; (f) Q. Shen, Y.-G. Huang, C. Liu, J.-C. Xiao, Q.-Y. Chen and Y. Guo, J. Fluorine Chem., 2015, 179, 14,
- 19 H. Li, X.-Y. Wang, B. Wei, L. Xu, W.-X. Zhang, J. Pei and Z. Xi, Nat. Commun., 2014, 5, 4508.
- 20 T. Seo, N. Toyoshima, K. Kubota and H. Ito, J. Am. Chem. Soc., 2021, 143, 6165.
- 21 R. Takahashi, A. Hu, P. Gao, Y. Gao, Y. Pang, T. Seo, S. Maeda, J. Jiang, H. Takaya, K. Kubota and H. Ito, Nat. Commun., 2021, 12, 6691.
- 22 (a) S. Maeda, K. Ohno and K. Morokuma, Phys. Chem. Chem. Phys., 2013, 15, 3683; (b) S. Maeda and Y. Harabuchi, Wiley Interdiscip. Rev.: Comput. Mol. Sci., 2021, 11, e1538.
- 23 For an example of the formation of organocalcium-based cubic structure, see: V. Knapp and G. Müller, Angew. Chem., Int. Ed., 2001, 40, 183.
- 24 For a review on calcium Lewis acid catalysis, see: J.-M. Begouin and M. Niggemann, Chem.-Eur. J., 2013, 19, 8030.